AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior version, and listings, of claims in the application:

1-13. (Cancelled)

- 14. (Currently amended) The method of claim 1, wherein prior to said culturing said oocyte is treated by A method of culturing an oocyte in vitro, comprising microinjecting into the cytoplasm of said oocyte a protective agent which (i) comprises a sugar, and (ii) is substantially non-permeating with respect to mammalian cell membranes and incubating said oocyte in a hypertonic medium having an osmolarity greater than 300 mosm.
- 15. (Previously presented) The method of claim 14, wherein said protective agent comprises at least one sugar selected from the group consisting of sucrose, trehalose, fructose, dextran, and raffinose.
- 16. (Previously presented) The method of claim 14, wherein said protective agent comprises at least one sugar selected from the group consisting of glucose, sorbitol, mannitol, lactose, maltose, and stachyose.

- 17. (Previously presented) The method of claim 14, wherein said protective agent comprises at least one sugar with a glass transition temperature greater than -50°C.
- 18. (Previously presented) The method of claim 17, wherein said protective agent comprises at least one sugar with a glass transition temperature greater than -30°C.
- 19. (Previously presented) The method of claim 14, wherein said protective agent comprises at least one sugar with a molecular weight greater than 120 daltons.
- 20. (Previously presented) The method of claim 14, wherein said protective agent comprises a glycolipid or a glycoprotein that comprises at least one sugar moiety derived from a sugar with a glass transition temperature greater than -50°C.
- 21. (Previously presented) The method of claim 14, wherein the cytoplasmic concentration of said sugar is less than or equal to about 1.0 M following microinjection.
- 22. (Previously presented) The method of claim 14, wherein the cytoplasmic concentration of said sugar is less than or equal to about 0.2 M following microinjection.
 - 23. (Currently amended) The method of claim 6, wherein prior to said culturing

said embryo is treated by A method of culturing an embryo in vitro comprising microinjecting into the cytoplasm of said embryo a protective agent which (i) comprises a sugar, and (ii) is substantially non-permeating with respect to mammalian cell membranes and incubating said embryo in a hypertonic medium having an osmolarity greater than 300 mosm.

- 24. (Previously presented) The method of claim 23, wherein said protective agent comprises at least one sugar selected from the group consisting of sucrose, trehalose, fructose, dextran, and raffinose.
- 25. (Previously presented) The method of claim 23, wherein said protective agent comprises at least one sugar selected from the group consisting of glucose, sorbitol, mannitol, lactose, maltose, and stachyose.
- 26. (Previously presented) The method of claim 23, wherein said protective agent comprises at least one sugar with a glass transition temperature greater than -50°C.
- 27. (Previously presented) The method of claim 23, wherein said protective agent comprises at least one sugar with a glass transition temperature greater than -30°C.

- 28. (Previously presented) The method of claim 23, wherein said protective agent comprises at least one sugar with a molecular weight greater than 120 daltons.
- 29. (Previously presented) The method of claim 23, wherein said protective agent comprises a glycolipid or a glycoprotein that comprises at least one sugar moiety derived from a sugar with a glass transition temperature greater than -50°C.
- 30. (Previously presented) The method of claim 23, wherein the cytoplasmic concentration of said sugar is less than or equal to about 1.0 M following microinjection.
- 31. (Previously presented) The method of claim 23, wherein the cytoplasmic concentration of said sugar is less than or equal to about 0.2 M following microinjection.
- 32. (New) The method of claim 14, wherein the osmolarity of said medium is greater than 320 mosm.
- 33. (New) The method of claim 32, wherein the osmolarity of said medium is greater than 340 mosm.
 - 34. (New) The method of claim 33, wherein the osmolartiy of said medium is

greater than 360 mosm.

- 35. (New) The method of claim 14, wherein said medium comprises a sugar selected from the group consisting of sucrose, trehalose, fructose, dextran, and raffinose.
- 36. (New) The method of claim 23, wherein the osmolarity of said medium is greater than 320 mosm.
- 37. (New) The method of claim 36, wherein the osmolarity of said medium is greater than 340 mosm.
- 38. (New) The method of claim 37, wherein the osmolartiy of said medium is greater than 360 mosm.
- 39. (New) The method of claim 23, wherein said medium comprises a sugar selected from the group consisting of sucrose, trehalose, fructose, dextran, and raffinose.